What is Claimed is:

- 1. An assay for the detection of Lyme disease infection comprising contacting a sample to be tested with a recombinant P37 FlaA protein antigen, incubating for sufficient time to allow formation of specific antibody-P37 FlaA protein antigen complexes, and detecting specifically bound antibody-P37/FlaA protein antigen complex.
- 2. An assay as in claim 1 wherein said recombinant P37/FlaA protein antigen has the amino acid sequence of amino acids 1 319 of the amino acid sequence of SEQ ID NO 2.
- 3. An assay as in claim 2 wherein said recombinant P37 protein antigen is expressed as a fusion protein with a fusion partner.
- 4. An assay as in claim 3 wherein said fusion protein partner is the approximately 38kDa T7 gene 10 product.
- 5. An assay as in claim 1 wherein said P37 protein antigen is immobilized on a solid support.
- 6. An assay as in claim 1 wherein said P37 protein antigen is derivatized with a detectable label.
- 7. An assay as in claim 1 wherein said antibody-P37 antigen complex is detected by specific protein binding to the antibody specific for P37.
- 8. An assay as in claim 1 wherein said detection uses chemiluminescent

cultures comprising constructing a DNA expression vector containing an expressible $1.1 \pm \Lambda_{\rm control}$ from DNA sequences transforming a soft able bases at the other soft forming and soft and the soft forming and the bases at the other soft forming and the bases are the other soft forming and the other soft forming are the other soft forming and the other soft forming are the other soft forming and the other soft forming are the other soft forming and the other soft forming are the other soft forming are the other soft forming are the other soft forming and the other soft forming are the other soft formin

vector; preparing large scale cell cultures from freshly transformed host cells, and not overnight cultures; inducing FlaA protein expression from said host cells in culture; and isolating recombinant FlaA protein.

- 10. A method as in claim 9 wherein the recombinant FlaA protein has the amino acid sequence of amino acids 1-319 of SEQ ID NO 2.
- 11. A method as in claim 10 wherein the recombinant FlaA protein is expressed as a fusion protein with a fusion partner.
- 12. A method as in claim 11 wherein the fusion partner is the approximately 38 kDa T7 gene 10 product.
 - 13. A method as in claim 5 wherein said host cell is an *E. coli* cell.
- 14. A recombinant FlaA produced using a method for producing recombinant FlaA protein from freshly transformed host cells comprising: constructing a DNA expression vector containing an expressible FlaA encoding DNA sequence; transforming a suitable host cell with said expression vector; plating out transformed host cells to generate individual fresh transformant colony of transformed host cells; preparing large scale primary cell cultures from transformed host cells taken from a fresh transformant colony, and not overnight cultures; allowing the primary cell culture to incubate for a period of time; inducing FlaA protein expression from said host cells in culture; and isolating recombinant FlaA protein.
- 15. A recombinant FlaA protein of claim 14, said protein having the amino

protein is expressed as a fusion protein.

- A recombinant FlaA protein as in claim 16 wherein the FlaA protein is expressed with a fusion partner that is the approximately 38 kDa T7 gene 10 product.
- 18. A recombinant FlaA protein as in claim 14 wherein said transformed host cell is an *E. coli* cell.